## AMENDMENTS TO THE CLAIMS

Claim 1 (currently amended). An article of manufacture comprising packaging material and a pharmaceutical composition contained within said packaging material, wherein said pharmaceutical composition is capable of modulating stimulating angiogenesis in a tissue associated with a disease condition, wherein said packaging material comprises a label which indicates that said pharmaceutical composition is administered to a patient for treating disease conditions by modulating stimulating angiogenesis, and wherein said pharmaceutical composition comprises consists essentially of at least about 0.1 weight percent of an isolated active or inactive Raf protein in a physiologically tolerable excipient or carrier therefore; wherein the active Raf protein is selected from the group consisting of c-Raf (SEQ ID NO: 2), a protein having the amino acid sequence corresponding to residues 306 through 648 of SEQ ID NO: 2, and Raf-caax (SEQ ID NO: 7); and the inactive Raf protein is selected from the group consisting of a protein having the amino acid sequence corresponding to SEQ ID NO: 2 having an amino acid other than lysine at residue 375, and a protein having the amino acid sequence corresponding to residues 1 through 305 of SEQ ID NO: 2.

Claim 2 (cancelled).

Claims 3 (currently amended). The article of manufacture of elaim 2 claim 1 wherein said active Raf protein is cRaf (SEQ ID NO: 2).

Claim 4 (cancelled).

Claim 5 (currently amended). The article of manufacture of claim 2 claim 1 wherein said active Raf protein is Raf-caax (SEQ ID NO: 7).

Claim 6 (currently amended). The article of manufacture of elaim 2 claim 1 wherein said tissue has poor circulation.

Claims 7-13 (cancelled).

Claim 14 (previously presented). The article of manufacture of claim 1 wherein said pharmaceutical composition is administered to a patient by intravenous, transdermal, intrasynovial, intramuscular, or oral administration.

Claim 15 (previously presented). The article of manufacture of claim 1 wherein said pharmaceutical composition is administered to a patient as a single dose intravenously.

Claims 16-40 (cancelled).

Claim 41 (currently amended). A pharmaceutical composition for stimulating angiogenesis in a target mammalian tissue comprising a therapeutic amount of an isolated consisting essentially of at least about 0.1 weight percent of an active Raf protein and in a pharmaceutically acceptable carrier or excipient, wherein the isolated active Raf protein is selected from the group consisting of c-Raf (SEQ ID NO: 2), a protein having the amino acid sequence corresponding to residues 306 through 648 of SEQ ID NO: 2, and Raf-caax (SEQ ID NO: 7).

Claims 42-67 (cancelled).